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Grey seal pup mortality is not explained by local breeding density on the island colony of North Rona, Scotland.

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Running head: Grey seal pup mortality

Abstract

Pre-weaning pup mortality in a UK grey seal breeding colony (North Rona) was examined in relation to the local adult female density over two breeding seasons. Accurate, fine scale maps of daily seal locations within a GIS database permitted the extraction of measures of female density determined at a range of spatial scales from 5m to 50m around each pup on the date of death. Adult densities around live pups provided the null distribution of adult densities which were tested against the distribution of densities where deaths occurred. Pre-weaning pup mortality at this colony was not related to any of our density measures. The proportion of pups dying each day showed no seasonal trends. Analysis of topographic features from a sub-metre Digital Terrain Model for locations where deaths occurred suggested that there were fewer mortalities in prime pupping habitats. This could have been a direct effect of habitat or the relative quality of mothers in these habitats. The distance separating mothers and pups tended to be greater on the periphery of the colony compared to more central areas. Pups in the peripheral areas may be more exposed to predation by gulls.

Key words: *Halichoerus grypus*, pinnipeds, breeding habitat, pup mortality, GIS

INTRODUCTION

Many species aggregate to breed. Advantages of group breeding include reduced risk from predation and an increased ability to detect and to exploit resources, including mates (Sutherland, 1996). However, these advantages are often traded against costs, such as increased risks of mortality, arising from density dependent processes of depletion, interference and predation (Sutherland, 1996). Density dependent mortality has been shown to occur in a wide range of vertebrates during their temporary breeding aggregations (reviewed by Sinclair, 1989), largely due to limitations on the availability of space or food. Much of this density dependent mortality impacts on juveniles (Sutherland, 1996).

Density dependent pup mortality has been suggested for some pinniped species at some breeding colonies (*Arctocephalus gazella*: Doidge, Croxall & Baker, 1984; *Arctocephalus australis*: Majluf, 1992; Harcourt, 1992; *Mirounga angustirostris*: Riedman, 1990; *Mirounga leonina*: Baldi *et al.*, 1996). However, comparisons of overall levels of mortality between different breeding colonies with measures of breeding density derived from colony-wide counts of animals do not consider an individual's environment at the specific location and time of its death (i.e. within a spatial and temporal scale at which the actions of conspecifics are likely to impact directly on an individual). Few studies have examined variation in mortality within a colony in relation to adult density and habitat variation measured at this 'local' resolution. Whilst some factors that may influence mortality are observed at larger scales (environmental variables such as weather conditions and tide), it can be argued that the impact of these factors on individuals depend upon their interaction with the individual at a local scale. For example, an individual's exposure to the prevailing

weather may depend upon the interaction of weather with local topographic features. Although weather parameters can easily be observed and measured at larger scales, they are experienced by individuals at local scales. Many other potential causes of individual deaths, particularly those associated with conspecifics (Baker, 1988; Baker & Baker, 1988) certainly operate at the scale of the individual and not the colony. Therefore, it is important to examine pup mortality with regard to these local scale measures. Similarly, as breeding densities often vary through a breeding season, it is necessary to assess local density at the time of death in order to ascertain if density is a potential cause of pup mortality.

Pinnipeds generally aggregate to breed (Bartholomew, 1970), frequently on island sites that are typically free from terrestrial predators. Many phocids rely upon stored energy reserves (blubber) to provision themselves and their pups during the annual breeding attempt. Thus, in species such as the grey seal (*Halichoerus grypus*) breeding in the UK, predation and local food resources are generally excluded from considerations of density dependent pup mortality within breeding aggregations. The primary candidate amongst potential determinants of pup mortality is likely to be interference from conspecifics. As such we would expect a relationship between local density and pup mortality. Density dependent pup mortality on grey seal colonies has been suggested by several studies (e.g. Boyd, Lockie & Hewer, 1962; Boyd & Campbell, 1971; Coulson & Hickling, 1964; Summers, Burton & Anderson, 1975). Breeding colony topography has also been implicated as a factor in pup mortality. Boyd *et al.* (1962), Coulson & Hickling (1964) and Summers *et al.* (1975) noted the higher mortality rates close to the major access routes to the colony. However, the relationship between topography and mortality has not been quantified at the scale of the individual.

Grey seals in Scottish waters aggregate to breed during each autumn, often on remote islands, such as North Rona. Individual females spend 18-20 days ashore, during which time they each bear and suckle one pup, come into oestrus towards the end of lactation and are mated. Of all the pups produced in a year, approximately 95% are born during a six week period (Harwood *et al.*, 1991; Hiby *et al.*, 1996). Hence, there is a turnover of females during the breeding season. At North Rona females generally remain with their pup throughout lactation (median daily movement of mothers is 10m from pupping location). Females aggregate around preferred habitat features (pools of water) (Twiss *et al.*, 2000a; Twiss, Thomas & Pomeroy, 2001), with variation in density related to habitat suitability (Twiss *et al.*, 2001). Males compete to maintain positions on the colony in areas that hold high densities of females (Anderson and Fedak, 1985; Twiss, 1991; Twiss, Anderson & Monaghan, 1998). Sex ratios on North Rona are typically in the order of 1male to seven females (Anderson and Fedak, 1985; Twiss *et al.*, 1998).

Here, we are able to investigate the occurrence of pre-weaning pup deaths within the grey seal breeding colony of North Rona using fine scale accurate mapping of pup and adult daily locations within a Geographical Information System (GIS). We use these data to test the null hypothesis that pup deaths are not related to measures of local adult female density. By linking these location data to a sub-meter accurate Digital Terrain Model (DTM) of the colony we also test the null hypothesis that levels of pup mortality are not related to local topography.

METHODS

Study site

The grey seal colony on the Scottish island of North Rona (59 06' N, 05 50' W) was established prior to the 1880s and has been the subject of studies of reproductive behaviour for 40 years (Boyd & Laws, 1962; Pomeroy *et al.*, 1999). The main seal breeding area is on the low lying northern peninsula (Fianuis), measuring 1.0x0.3 km. Our study area encompasses the southern portion of this peninsula covering 185000m². The duration of the study period in both the 1997 and 1998 breeding seasons extended from the onset of breeding at North Rona to past the time of maximum number of adults ashore (see Fig. 1a and b).

Mapping of seal locations

During the 1997 and 1998 breeding seasons the daily locations of all seals within the North Rona study area were mapped precisely on detailed fine grain, geo-referenced maps of the breeding colony. Maps incorporating detailed habitat features and a 10x10m grid were derived from digitised geo-rectified high resolution aerial photographs (Twiss *et al.*, 2000a, 2000b, 2001). Every seal was identified by sex and age and individual identities where known, with all pups being classified into either age categories 'I' to 'V' (based on the well established descriptive age classes of Boyd *et al.*, 1962) or as dead pups. All these daily maps and classifications were digitised into an ARC-INFO GIS database (Twiss *et al.*, 2000 a, 2000b, 2001).

GIS data extraction

From these GIS coverages (topologically linked geographic features with their associated descriptive data) of seal locations for each breeding season we extracted the locations of all dead pups and their associated dates of observation into a new GIS

coverage. Most dead pups remain on the North Rona breeding colony for a considerable period (see results). Therefore, in our data-set successive daily maps contained repeat records of the same dead pup. These repeat records of dead pups formed readily distinguishable clusters of locations within our composite GIS coverage of dead pups (dead pups did move marginally; artefactually through minor mapping errors, but also in reality through the actions of other seals and in particular scavenging gulls). Hence, for each cluster we selected the location with the earliest date, and used this as our representative location for the mortality and the date on which it was first recorded as its day of death. These selected 'first' occurrences of each dead pup were generated into a new coverage, one for each breeding season. We then used these coverages to automatically extract from daily coverages of female locations the number of adult females within a specified radius of each dead pup on the specific date on which the pup died. This process was repeated to extract the number of adult females in successively increasing radii, from 5m to 50m in 5m increments. Thus, this process provides measures of the local adult female density, on a range of scales, around the location of each dead pup on the day of its death. This day-specific approach combined with the flexibility in the scale of density measurement, provides a more realistic assessment of potential density effects within the colony, than, for example, comparisons of the number of dead pups in sub sections of the colony with seasonal averages or maximum observed adult densities. Furthermore, it provides the capability to detect relationships that may operate at scales beyond the dead pups immediate vicinity, for example, if abandoned or starveling pups wander away from their natal sites prior to death.

Estimation of the percentage pup mortality

The total number of first occurrences of dead pups provides a count of the number of dead pups over each season within our study site. To convert this to a percentage of the total number of pups born in the study area, we extracted from our coverages of pup locations the number of pups of stage I to IV (stage V pups are fully moulted and usually weaned pups). We plotted the daily number of these pups and computed the area under this graph (the sum of daily numbers i.e. total number of pup-days) for each season. This sum was then divided by 18, the average duration from birth to weaning. To this resulting value we added the number of dead pups to provide an estimate of the total number of pups born in the study area. From this we generated an overall estimate for the seasonal percentage of mortality.

Temporal sequence of mortalities

The seasonal change in the rate of mortalities was examined with respect to both the numbers of adult females and the numbers of all pups on a daily basis.

Local adult female density and pup mortality

In order to test whether mortalities tended to occur more often than expected in areas of higher adult densities it was necessary to provide null distributions of the 'available' adult densities on the colony. Using a similar GIS based process to that described above, we extracted the densities of females (for each of the 5m to 50m radii) around every live pup's location on every day for each season. Inevitably, the density of females around any one pup on any one date was unlikely to be independent of the density on the preceding or following days. However, these records numbered 5835 for 1997 and 5848 for 1998. Therefore, we generated

probability density functions for our null distributions by repeating 10000 bootstrapping procedures in which we resampled with replacement 1% of these records. This process was repeated for each spatial scale and for year of the two breeding seasons. Seals do not utilise all the area mapped within our GIS but prefer specific habitat features (Twiss *et al.*, 2000a, 2001). Therefore, the use of live pup locations to sample adult densities provided a more appropriate measure of the available densities than would be achieved by random locations taken throughout the study area which would inevitably include samples from parts of the colony devoid of seals.

For each season and each sampling radii of adult female density we compared the distributions of adult densities around mortalities with the corresponding distribution derived from our bootstrapping of densities around live pups. Comparisons were made by Kolmogorov-Smirnov 2 sample tests.

Habitat and mortality

A sub-meter accurate DTM of the study area (Mills, Newton & Twiss, 1997; Twiss *et al.*, 2000a, 2000b, 2001) provides accurate descriptions of the physical habitat for any location. These topographic measures include;

- (1) elevation above mean sea level (m)
- (2) Slope (degrees)
- (3) "Cost distance" to "nearest" pool of water,
- (4) "Cost distance" to "nearest" access point from the sea.

The cost–distance variables provide an index for each location of the relative ease of moving over an irregular terrain from that location to the nearest pool or access point, though nearest is defined as the easiest to get to (least cost). Large physical features, such as cliffs present impenetrable barriers to movement in these models, requiring circumnavigation and hence longer routes and higher costs. Differing slopes also present differing costs to movement. These models provide an accurate representation of the relative ease of locomotion across the colony, and have been used to demonstrate seals’ preferences for breeding close to access points and / or close to pools of water (see Twiss *et al.*, 2000a, 2000b, 2001 for further details of cost-distance modeling procedures).

Each of these variables was selected for the 2m grid square in which each dead pup was found. Null distributions of ‘available’ habitat were generated in the same fashion as for the adult densities above; using 10000 bootstrapping procedures resampling with replacement 1% of the habitat values associated with live pup locations. We then compared the distribution of habitat variables for dead and live pups.

Female-pup proximity relative to position in the colony

We hypothesised that females on the periphery of the colony, in lower density areas may not attend their pups as closely as those in denser, central areas. A useful measure of centrality of location in the colony is provided by our index of cost-distance to access (Twiss *et al.*, 2001, Pomeroy *et al.*, 2001.). Females prefer to pup at an intermediate distance from access points, as access points tend to experience more traffic, exposure to waves and increased numbers of transient males (Twiss *et al.*, 2001). Using coverages of all live pups and adult females we determined the

distance between every pup on each day and their nearest adult females using GIS proximity functions (Twiss *et al.*, 2000a). We compared these distances between pup and nearest female for categories of cost-distance to access (divided into bins of 50 units in an index ranging from 0 to 300) using Kruskal-Wallis tests. A similar analysis was conducted to examine the pup to nearest female distances in relation to cost-distance to pools of water (divided into bins of five units in an index ranging from 0 to 35).

RESULTS

Persistence of corpses on the colony

Dead pups on North Rona remained visible, and were therefore mapped, for an average of 9.0 ± 1.0 consecutive days in 1997 and 10.2 ± 1.2 days in 1998. However, many of the dead pups (32% in 1997 and 35% in 1998) remained visible up to and including our last observation date, therefore, the above values represent underestimates of the true persistence of corpses on the colony.

Estimates of the percentage pup mortality

Estimated pup mortality rates were 14.6% (52/357) in 1997 and 14.4% (51/354) in 1998.

Temporal sequence of mortalities

Fig. 1a and b shows the change in number of pups, adult females and dead pups during the 1997 and 1998 seasons respectively. The number of dead pups is

represented as both the total number of corpses recorded on each date (which approximates to cumulative deaths) and as the number of deaths (first occurrences of dead pups) per day (daily mortality rate). Note that the numbers of corpses and deaths have been exaggerated by a factor of ten for clarity in these figures. Table 1 presents the correlations of these two daily measures of mortality with date during the breeding season, the total number of adult females present per day and thirdly, the total number of pups present per day. The total number of corpses recorded on each date correlates highly with both the number of females and the number of pups present (as the number of corpses is a cumulative measure, no correlation was attempted with date). However, significant positive correlations also existed between the daily mortality rate and either date, the number of females and the number of pups per day. Conversely, when the daily mortality rate is expressed as a percentage of the total number of pups present per day then no significant correlations remained.

Local adult female density and pup mortality

In both breeding seasons, the frequency distributions of densities of adult females around pup mortalities showed no significant differences from the distributions of available female densities at $p < 0.05$ for all spatial scales at which female densities were sampled with the exception of the smallest, 5m, scale (Table 2). Examination of the cumulative frequency distributions for dead and live pups for the 5m spatial scale (Fig. 2a and b) suggests that dead pups tended to occur more frequently than live pups in locations where there were no or few adult females within 5m of the dead pup (i.e. at the lowest density categories).

Habitat and mortality

Our analysis of the nature of the topography at locations where deaths occurred compared to the null distribution of topographies at live pup locations provided inconsistent results between years. In the 1998 season there were no significant differences at $p < 0.01$ between the topography at dead and live pup locations (Table 3). However, in 1997, the frequency distributions of both elevation and cost distance to access differed for dead and live pups (Table 3). Examination of the cumulative frequency distributions of elevation values for dead and live pups (Fig. 3) indicated that dead pups occurred more frequently than live pups at the lowest elevations and less frequently at intermediate elevations (approximately 10-25m). In terms of cost-distance to access (Fig. 4), dead pups occurred more frequently in the immediate vicinity of access points (low cost-distance indices) and less frequently than live pups at moderate indices for cost-distance to access (indices of approximately 60 to 100).

Female-pup proximity relative to position in the colony

The distances separating pups and their nearest adult female tended to (i) increase with distance from access points (Kruskal-Wallis tests: 1997 $\chi^2 = 98.7$, d.f. = 5, $p < 0.001$, 1998 $\chi^2 = 53.2$, d.f. = 5, $p < 0.001$) and (ii) increase with distance from pools of water (Kruskal-Wallis tests: 1997 $\chi^2 = 142$, d.f. = 6, $p < 0.001$, 1998 $\chi^2 = 47.6$, d.f. = 6, $p < 0.001$).

DISCUSSION

Contrary to earlier studies which examined pup mortality rates in relation to colony wide measures of pup production (Boyd *et al.*, 1962; Boyd & Campbell, 1971; Coulson & Hickling, 1964; Summers *et al.*, 1975), our fine scale analyses failed to demonstrate any evidence that pre-weaning pup mortality increased with adult densities at this breeding colony. The densities in which mortalities occurred encompassed the full range of available adult densities on the colony. The occurrence of pup deaths was not related to local density of adult females irrespective of the scale at which local density was measured with the exception of our smallest scale of analysis. At this 5m scale our analysis again revealed no indication of mortality increasing with local density. In fact, dead pups were found more frequently in locations with no adult females within 5m.

Our analyses were repeated at a range of scales that extended from the pup's immediate vicinity to well beyond any distance at which adult female density could possibly directly influence individual pup deaths. The significant positive correlation between the daily mortality rate and the number of females per day may give a misleading appearance of a relationship between density and mortality when viewed at this relatively coarse scale. It is likely that this relationship is mediated through a mutual correlation with date. This was supported by the fact that there was no correlation when the mortality rate was expressed as a percentage of the daily number of pups. Thus, the absolute number of deaths increased with increasing female numbers, but the proportion of deaths remained constant as increasing numbers of pups were born. Furthermore, our fine scale analysis of local adult female density and pup deaths refuted any notion that mortalities increased with local adult female density within the range of densities occurring in the North Rona colony. Animal densities on grey seal breeding colonies tend to be lower than those seen in colonies

of elephant seals (Riedman, 1990; Baldi *et al.*, 1996) and of many otariid species (Harcourt, 1992; Riedman, 1990) in which pup mortality is density dependent. Even amongst the most dense aggregations, female grey seals maintain a distance of approximately 1.5 body lengths between themselves and their neighbours (Twiss *et al.*, 2000a). Our results suggest that local adult densities on the day of death do not directly cause pup deaths. It is possible that more complex, compound effects of local density on a number of days preceding death may provide a link between adult density and death. Abandoned or starveling pups may wander from natal sites. Our finding that dead pups tended to occur more frequently in locations where no potential mothers were found within 5m radii may reflect either this, or that mothers leave a dead pup (although mothers have been observed maintaining close contact with dead pups for a number of days, pers. obs. PPP, SDT). A further consideration is the effect of adult males. Whilst densities of males, given the skewed sex ratio, are very low compared to adult females, it is possible that pup mortalities, or at least injuries that subsequently lead to death, result as a by-product of male-male aggression. It is clearly necessary to quantify these parameters, and future studies should endeavour to; (a) track pup locations in relation to local density prior to death, (b) quantify the movements of live pups once they have been abandoned, (c) quantify maternal attendance, in terms of proximity to pups, in relation to local density (d) quantify local male densities and more importantly the frequency and intensity of inter-male aggression with respect to location.

We detected no seasonal trend in the rate of pup mortality. This indicates that for the period of these studies, there is no cost in terms of increased pup loss to later breeding females. Boyd & Campbell (1971) stated that an accurate measure of seasonal trends in daily mortality rates would aid in estimating pup production, but

stated that daily ground counts would incur too much disturbance (Summers *et al.*, 1975 provided daily mortality rates at weekly intervals). Here, we provide such data, on a daily basis, with no disturbance, and demonstrate that daily mortality rates are proportional to the number of pups present in any day.

Our estimates of overall seasonal mortality rates of approximately 14.5% matched closely estimates derived from more direct means (ground counts) at both this colony and others. Mortality estimates for the southern part of the Fianuis peninsula of North Rona provided by Boyd & Laws (1962) were 15% of 542 births and 13% of 510 births in 1959 and 1960 respectively, whilst Boyd & Campbell (1971) provide estimates ranging from 10 to 20% over the years 1962 to 1968 with an average of 14.5%. The study area used in these studies is larger than ours but encompasses our study area. Summers *et al.* (1975), studying the same area as we use here, estimated 30% mortality from 393 births in the 1972 season. Further afield, Baker & Baker (1988) estimated mortality rates for the Isle of May (Scotland) ranging from 13.0 to 15.7%, Coulson and Hickling (1964) presented an average mortality rate of 14.9% for the Farne Islands (England), and Davies (1949) recorded 15% for beach breeding seals in Pembrokeshire (Wales). The concurrence of these estimates suggests that our approach provides a good measure of the number of dead pups. Furthermore, the remarkable consistency of most of these estimates of mortality rates, despite the fact that the various colonies mentioned above have differing topographies and seal densities to North Rona (Pomeroy, Twiss & Duck, 2000; Twiss *et al.*, 2000a, 2001), might suggest that mortality rates are linked to larger scale effects. It may be that the mortality rate reflects the proportion of breeding females that have insufficient energy to successfully rear a pup in any one year, and so be related to factors outside the breeding season that affect female body condition (although if this was the case one

might expect more inter-annual variation in mortality rates). If such energetic considerations are responsible for underlying mortality rates, then it is possible that this might mask more subtle density-dependent effects. Subsequent studies of local density effects on mortality should endeavour to control for maternal condition.

Density dependent mortality in breeding aggregations may be absent where advantages accrue to centrally located individuals and yet central locations attract the highest densities of individuals (Sutherland, 1996). Competition for desirable central, high density locations may lead to occupancy of these prime sites by higher quality individuals. These individuals may then offset the potential density dependent effects by their better parental qualities and through their enhanced location. Amongst grey seals, female pupping site preferences in relation to variation in habitat quality at UK grey seal breeding colonies (Pomeroy, Twiss & Duck, 2000; Twiss *et al.*, 2000a, 2001) has been linked with differences in female quality (Pomeroy *et al.*, 1999, 2001). Therefore, individual quality could confound density dependent effects in grey seal colonies such as North Rona.

Our analyses indicated a possible role for habitat, at least in the 1997 breeding season. Increased mortality in the immediate vicinity of access points is expected (Baker, 1988; Pomeroy *et al.*, 1994), as these areas form major transit routes for adult seals, leading to increased disturbance of mothers and pups (Boyd *et al.* 1962, Coulson & Hickling 1964, Summers *et al.* 1975). Our results from 1007 suggested such an effect, with increased mortality at low elevations and cost-distances to access (effectively in, or around, access points). Twiss *et al.* (2000a, 2001) demonstrated that locations with intermediate distances to access points (i.e. not being immediately in the access points) and/or close proximity to pools of water are preferred pupping locations for breeding grey seals. Likewise, Twiss *et al.* (2000a) provided evidence

that local topography can determine the level of behavioural investment in pups by mothers. Pomeroy *et al.* (2001) extended this idea and provided evidence of high pup growth rates and potentially increased genetic success for mothers occupying these prime habitats. The result presented here from the 1997 season indicate an apparent reduction in the number of dead pups occurring at these same intermediate elevations and cost-distance to access indices (Fig. 4). Thus, reduced pup mortality in prime habitats may to be a direct consequence of habitat quality. However, since these patterns were not observed in both the breeding season examined here, it is perhaps more likely to be effected indirectly via the increased quality of animals occupying these sites. Linking long-term studies of individual females with habitat, local density and detailed behavioural observation should enable the causal direction of this relationship to be determined.

Finally, it may be necessary to reconsider the role of predation. Whilst there are no terrestrial predators of grey seal pups on North Rona, Greater and Lesser Black-Back gulls attack weakened or unprotected pups (pers. obs. PPP, SDT). Again, both maternal quality and habitat may be important here. Many otariids make large daily movements whilst breeding, often in order to access pools or the sea for thermoregulation (Harcourt, 1992). Grey seals at North Rona appear to do the same (Redman, Pomeroy & Twiss, 2001; Twiss *et al.*, 2000a, 2001, in press), but generally do not have to move too far from their pup due to the abundance of pools. However, pups born further from pools appear to spend more time further away from their mothers as their mothers spend longer commuting to pools. This may make these pups more vulnerable to predation by gulls. Thus, predation risk is likely to be a function of habitat (pupping site) quality, and potentially individual quality. Further observations will be required to address this question.

Whilst the pattern of pup mortalities on North Rona are not related to local variation in adult female density, our results suggest a possible role for both habitat and maternal quality in reducing pup deaths. These results have implications for studies of both the behavioural ecology of grey seals and also for applied models of population growth that require estimates of pup mortality on breeding colonies. Our results also illustrate the importance of examining individual based processes at the appropriate spatial scale.

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Table 1: Spearman's rank correlations (R) of date, daily number of adult females and daily number of live pups with: (a) the number of corpses recorded per day (as the number of corpses is effectively a cumulative measure, no correlation was attempted with date), (b) the number of deaths per day (daily mortality rate) and (c) daily mortality rate expressed as a percentage of total number of pups (live and dead) present per day. Significant (2-tailed) correlations are emphasised in bold.

		(a) Number of corpses observed per day (cumulative deaths)		(b) Number of deaths per day (daily mortality rate)		(c) Daily mortality rate as a percentage of the total number of pups present on each day	
		1997 (n = 27)	1998 (n = 28)	1997 (n = 27)	1998 (n = 28)	1997 (n = 27)	1998 (n = 28)
Date	R	-	-	0.49	0.63	-0.27	0.04
	p			0.001	<0.001	0.17	0.86
Number of adult females	R	0.92	0.92	0.49	0.56	-0.24	-0.08
	p	<0.001	<0.001	0.009	0.002	0.24	0.97
Number of live pups	R	0.97	0.97	0.47	0.63	-0.30	0.03
	p	<0.001	<0.001	0.014	<0.001	0.14	0.87

Table 2. Results of Kolmogorov- Smirnov (KS) tests of the frequency distribution of female densities around pup mortalities on the day of death compared against the null distribution from 10000 resamplings of adult densities around live pup locations. Analysis repeated using female densities determined for a range of radii around each pup location, from 50m to 5m. Sample sizes are 52 dead pups and 578144 live for 1997 and 51 dead and 578796 live in 1998. Significance difference at $p < 0.01$ are emphasised in bold.

Year		Radius (m) around pups in which adult female density was determined									
		50	45	40	35	30	25	20	15	10	5
1997	KS Z	1.23	1.31	1.23	1.08	1.31	1.03	0.78	0.57	0.93	1.64
	P	0.10	0.06	0.1	0.19	0.07	0.24	0.57	0.91	0.35	0.009
1998	KS Z	1.20	1.09	1.10	1.05	1.13	0.89	1.00	1.34	0.99	2.53
	P	0.11	0.19	0.18	0.22	0.16	0.40	0.27	0.06	0.28	<0.001

Table 3: Results of Kolmogorov- Smirnov (KS) tests of the frequency distribution of four topographic variables derived from the locations provided by dead pups and from 10000 resamplings of live pup locations. Sample sizes are 52 dead pups and 578144 live for 1997 and 50 dead and 572974 live in 1998. Significance difference at $p < 0.01$ are emphasised in bold.

Year		Topographic parameter			
		elevation	slope	Cost-distance to access	Cost-distance to pool
1997	KS Z	1.67	1.28	1.79	0.84
	P	0.008	0.08	0.003	0.49
1998	KS Z	0.72	1.59	0.82	1.51
	P	0.68	0.013	0.52	0.021

Figure Legends:

Fig. 1a & b: Plots of the daily numbers of adults females, live pups, pup corpses (cumulative number of deaths) and pup deaths (daily mortality rate) within the North Rona study area during the (a) 1997 and (b) 1998 breeding seasons. *NB: Numbers of pup corpses and pup deaths have been multiplied by a factor of 10 to give a clearer graphical presentation of changes in relative number with date.

Fig. 2a & b: Cumulative frequency histograms of densities of adult females within 5m of dead pups and live pups during the (a) 1997 and (b) 1998 breeding seasons.

Fig. 3: Cumulative frequency histogram of the elevation values at locations where dead pups and live pups occurred during the 1997 breeding season.

Fig 4: Cumulative frequency histogram of the cost-distance to access values at locations where dead pups and live pups occurred during the 1997 breeding season.

Fig. 1a: Twiss, Duck & Pomeroy: Grey seal pup mortality.

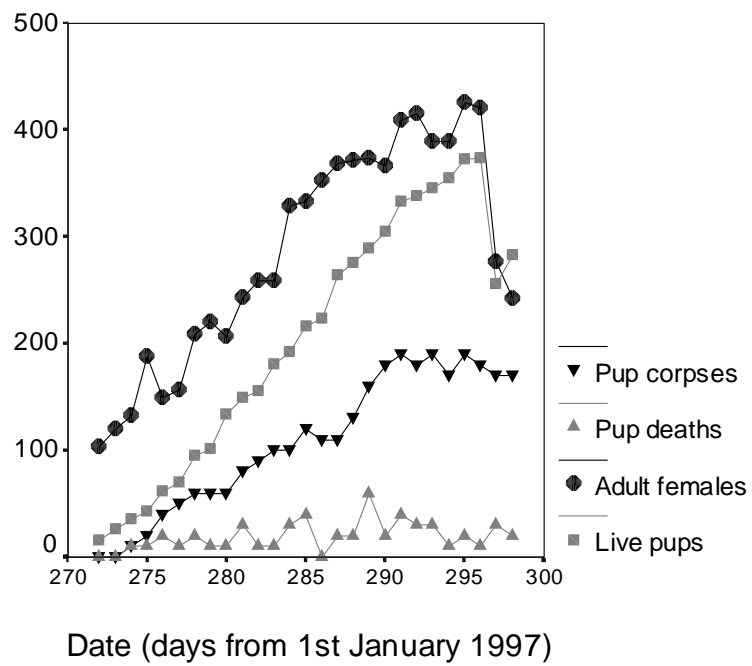


Fig. 1b: Twiss, Duck & Pomeroy: Grey seal pup mortality.

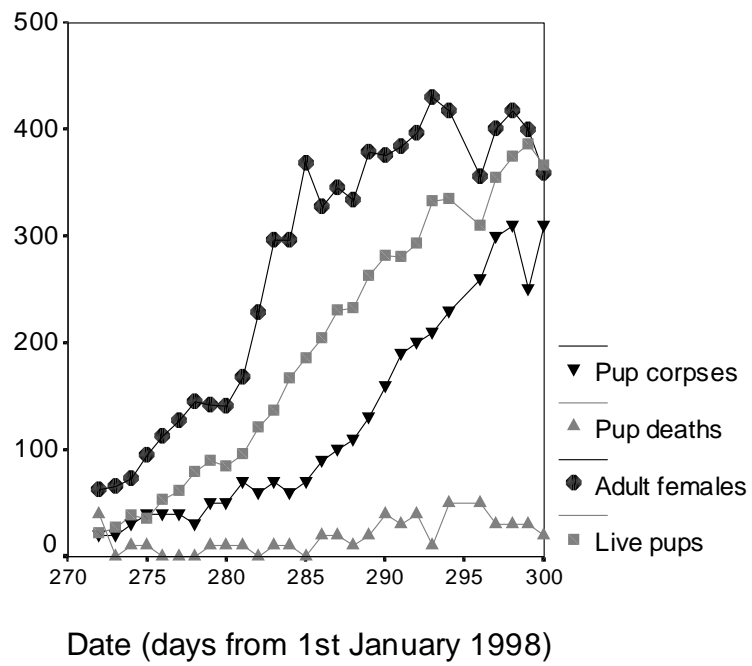


Fig. 2a: Twiss, Duck & Pomeroy: Grey seal pup mortality.

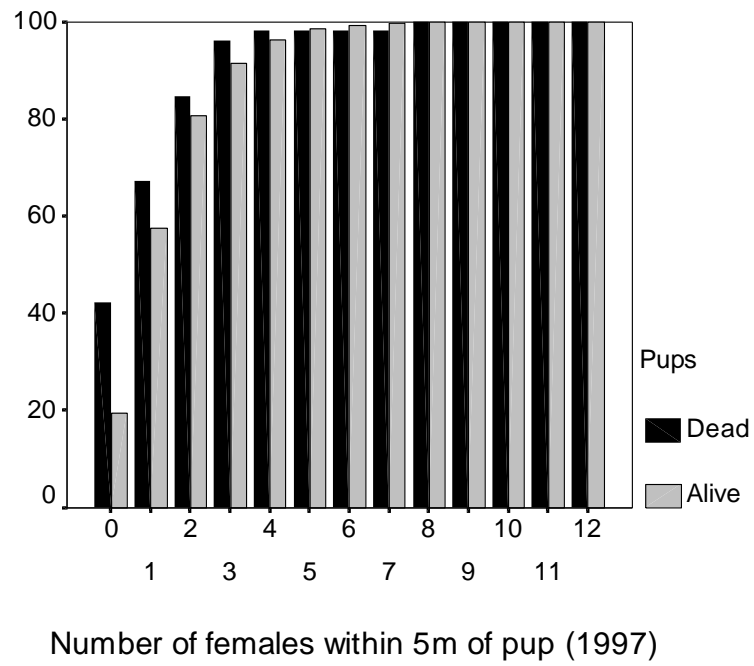


Fig. 2b: Twiss, Duck & Pomeroy: Grey seal pup mortality.

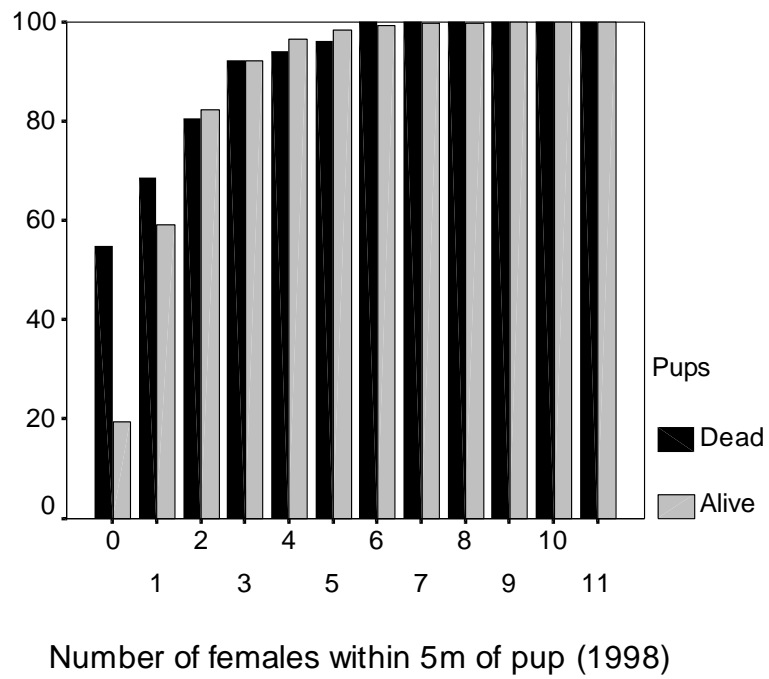


Fig. 3: Twiss, Duck & Pomeroy: Grey seal pup mortality.

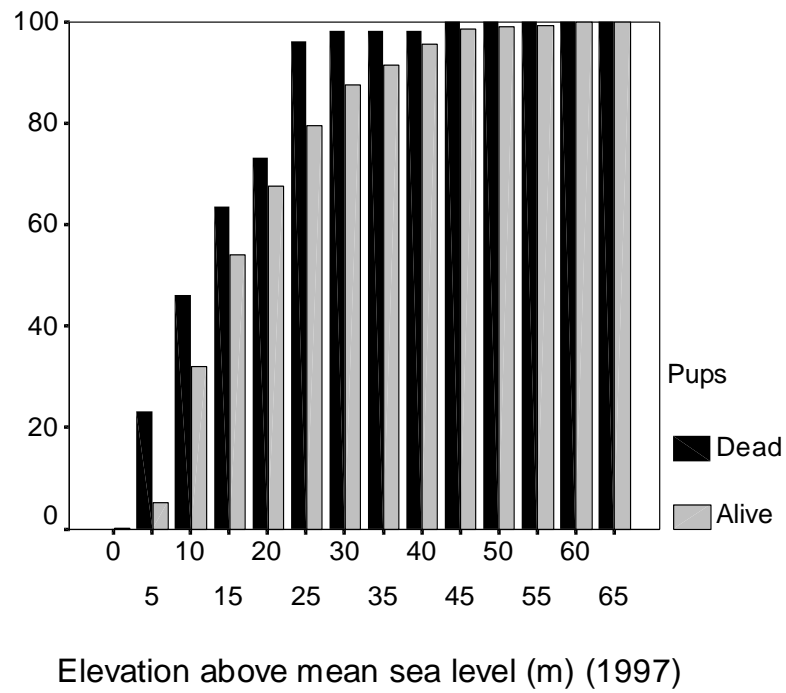


Fig. 4: Twiss, Duck & Pomeroy: Grey seal pup mortality.

